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Life Sciences Reporting Summary

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Please do not complete any field with "not applicable" or n/a. Refer to the help text for what text to use if an item is not relevant to your study. For final submission: please carefully check your responses for accuracy; you will not be able to make changes later.

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1.	Sami	ole	size

Describe how sample size was determined.

We used the 1000 Genomes Project whole genome samples where there is 26 populations and each population have on average 100 individuals. Therefore for simulation we chose 100 individuals (200 haplotypes).

2. Data exclusions

Describe any data exclusions.

No data were excluded.

3. Replication

Describe the measures taken to verify the reproducibility of the experimental findings.

We have done extensive simulations with different parameters that are explained in details in the Supplementary information. We also tested the performance of the method in well-characterized examples of human population.

4. Randomization

Describe how samples/organisms/participants were allocated into experimental groups.

As provided by 1000 Genomes Project, samples are grouped by population. For example, British in England and Scotland (GBR), Yoruba in Ibadan, Nigeria (YRI), and so forth.

5. Blinding

Describe whether the investigators were blinded to group allocation during data collection and/or analysis.

Data is provided by 1000 Genomes Project and we didn't change anything regarding the data.

Note: all in vivo studies must report how sample size was determined and whether blinding and randomization were used.

6. Statistical parameters

For all figures and tables that use statistical methods, confirm that the following items are present in relevant figure legends (or in the Methods section if additional space is needed).

n/a	Confirmed
	The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement (animals, litters, cultures, etc.)
	A description of how samples were collected, noting whether measurements were taken from distinct samples or whether the same sample was measured repeatedly

| | | | | A statement indicating how many times each experiment was replicated

- The statistical test(s) used and whether they are one- or two-sided

 Only common tests should be described solely by name; describe more complex techniques in the Methods section.
- | | | | | | A description of any assumptions or corrections, such as an adjustment for multiple comparisons

Test values indicating whether an effect is present

Provide confidence intervals or give results of significance tests (e.g. P values) as exact values whenever appropriate and with effect sizes noted.

A clear description of statistics including central tendency (e.g. median, mean) and variation (e.g. standard deviation, interquartile range)

Clearly defined error bars in <u>all</u> relevant figure captions (with explicit mention of central tendency and variation)

See the web collection on statistics for biologists for further resources and guidance.

Software

Policy information about availability of computer code

7. Software

Describe the software used to analyze the data in this study.

selscan (v1.1.0a); Szpiech et al. (2014); https://github.com/szpiech/selscan SCCT (v1.1); Wang et al. (2014); https://github.com/wavefancy/scct CMS (v2.0); Grossman et al. (2010); https://github.com/broadinstitute/cms

For manuscripts utilizing custom algorithms or software that are central to the paper but not yet described in the published literature, software must be made available to editors and reviewers upon request. We strongly encourage code deposition in a community repository (e.g. GitHub). *Nature Methods* guidance for providing algorithms and software for publication provides further information on this topic.

Materials and reagents

Policy information about availability of materials

8. Materials availability

Indicate whether there are restrictions on availability of unique materials or if these materials are only available for distribution by a third party.

no unique material were used.

9. Antibodies

Describe the antibodies used and how they were validated for use in the system under study (i.e. assay and species).

no antibodies were used.

10. Eukaryotic cell lines

- a. State the source of each eukaryotic cell line used.
- b. Describe the method of cell line authentication used.
- c. Report whether the cell lines were tested for mycoplasma contamination.
- d. If any of the cell lines used are listed in the database of commonly misidentified cell lines maintained by ICLAC, provide a scientific rationale for their use.

no eukaryotic cell line were used

▶ Animals and human research participants

Policy information about studies involving animals; when reporting animal research, follow the ARRIVE guidelines

11. Description of research animals

Provide all relevant details on animals and/or animal-derived materials used in the study.

no animals were used

Policy information about studies involving human research participants

12. Description of human research participants

Describe the covariate-relevant population characteristics of the human research participants.

The study didn't involve human research participants